Amendments to the Claims:

This listing of claims will replace all prior versions and listing of claims in the application.

Claims 4-10, 24-25 and 28-44 are amended.

Listing of Claims:

1. (Original) A compound of general formula:

Wherein:

 R_1 is selected from H and CH_3 , and R_2 is selected from H and OH, or R_1 and R_2 together form an optionally substituted phenyl ring which is fused to the pyridine ring; and

R₃ is selected from H, CH₃, CH₂0H and

R₄ is selected from H, CH₃, CH₂OH,

R₅ is selected from H, phenyl, halogen-substituted phenyl and

Wherein R_6 and R_7 are each independently selected from H, Na^+ , K^+ , alkyl and optionally substituted aryl, and X and Yare each independently selected from H, OH and F, or at least one of X and Y is an heteroatom and together with R_3 forms a bridge with the proviso that R_4 is

$$X$$
 Y
 O
 OR_6
 OR_7
 OR_7

and N-oxides thereof, and biologically acceptable salts thereof.

- 2. (Original) The compound according to claim 1, wherein said halogen-substituted phenyl is a fluoro-substituted phenyl.
- 3. (Original) The compound according to claim 1, wherein said halogen-substituted phenyl is $p-C_6H_4F$.
- 4. (Currently Amended) The compound according to any one of claims 1 to 3 claim 1, wherein said heteroatom is selected from 0 and S.
- 5. (Currently Amended) The compound according to any one of claims 1 to 3 claim 1, wherein said heteroatom is O.
- 6. (Currently Amended) The compound according to any one of claims 1 to 5 claim 1, wherein said bridge is selected from -CH2-, -CH₂CH₂- and -CH₂CH₂-.
- 7. (Currently Amended) The compound according to any one of claims 1 to 5 claim 1, wherein said bridge is a methylene bridge.
- 8. (Currently Amended) The compound according to any one of claims 1 to 7 claim 1, wherein said alkyl is a C_1 to C_6 straight or branched alkyl.
- 9. (Currently Amended) The compound according to any one of claims 1 to 7 claim 1, wherein said alkyl is t-butyl.
- 10. (Currently Amended) The compound according to any one of claims 1 to 9 claim 1, wherein said aryl is phenyl or naphthyl.
- 11. (Original) The compound according to claim 1, wherein R_1 , R_2 , R_3 , R_5 are all H and R_3 is

12. (Original) The compound according to claim 1, wherein R_1 , R_2 , R_3 , R_5 are all H and R_4 is

13. (Original) The compound according to claim 1, wherein R_1 , R_2 , R_3 , R_4 are all H and R_5 is

14. (Original) The compound according to claim 1 wherein R_1 and R_2 together form an optionally substituted phenyl ring which is fused to the pyridine ring; R_3 and R_5 are both H; and R_4 is

15. (Original) The compound according to claim 1, wherein R_1 and R_3 are both CH_3 ; R_2 is OH; R_5 is H; and R_4 is

16. (Original) The compound according to claim 1, wherein R_1 and R_4 are both CH_3 ; R_2 is OH; R_5 is H; and R_3 is

17. (Original) The compound according to claim 1, wherein R_1 is CH_3 ; R_2 is OH; R_3 is CH_2OH ; R_5 is H; and R_4 is

18. (Original) The compound according to claim 1, wherein R_1 is CH_3 ; R_2 is OH; R_3 is CH_2OH ; R_5 is H; and R_4 is

$$X$$
 Y
 O
 P
 OR_7

19. (Original) The compound according to claim 1, wherein R_1 is CH_3 ; R_2 is OH; R_5 is CH_2OH ; R_5 is C_6H_5 ; and R_4 is

20. (Original) The compound of claim 1, wherein R_1 is CH_3 ; R_2 is OH; R_3 is CH_2OH ; R_5 is $p-C_6H_4F$; and R_4 is

- 21. (Original) A compound according to claim 1, wherein R_5 is $P\text{-}C_6H_4F$.
- 22. (Original) A compound according to claim 1 selected from: [Hydroxy-(5-hydroxy-4-hydroxymethyl-6-methyl-2-phenyl-pyridin-3-yl)-methyl]-phosphonic acid; {[2-(4Fluoro-phenyl)-5-hydroxy-4-hydroxymethyl-6-methyl-pyridin-3-yl]-hydroxymethyl}phosphonic acid; [Hydroxy-(4-pyridin-2-yl-phenyl)-methyl]-phosphonic acid; [Fluoro(4-pyridin-2-yl-phenyl)-methyl]-phosphonic acid; (Hydroxy-quinolin-3-yl-methyl)phosphonic acid; (Fluoro-quinolin-3-yl-methyl)-phosphonic acid; [Hydroxy-(5hydroxy-4, 6-dimethyl-pyridin-3-yl)-methyl]-phosphonic acid; (Hydroxy-pyridin-4-yl-methyl)-phosphonic acid; (Hydroxy-pyridin-4-yl-methyl)-phosphonic acid; (Hydroxy-6-methyl-1 ,3-dihydro-furo[3,4-c]pyridin-3-yl)-phosphonic acid; [(3,7Dihydroxy-6-methyl-1 ,3-dihyrdo-furo[3 ,4-c]pyrid i n-3-yl)-difluoromethyl]-phosphonic acid; and nicotinyl phosphonates thereof, N-oxides thereof, phosphonate esters thereof and biologically acceptable salts thereof.

23. (Original) A compound according to claim 1 comprising:

- 24. (Currently Amended) A pharmaceutical composition comprising a therapeutically effective amount of a compound according to anyone of claims 1 to 23 claim 1 and a pharmaceutically acceptable carrier.
- 25. (Currently Amended) The compound according to anyone of claims 1 to 24 claim 1, wherein at least one polar group is blocked by a lipophilic moiety capable of being enzymatically cleaved off after absorption into the circulatory system.
- 26. (Original) The compound according to claim 25, wherein said lipophilic moiety is an ester.
- 27. (Original) The compound according to claim 25, wherein said lipophilic moiety is a phosphonate ester.
- 28. (Currently Amended) A method of treating hypertension in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to anyone of claims 1 to 27 claim 1 in a unit dosage form.
- 29. (Currently Amended) A method of treating myocardial infraction in a mammal

comprising administering to the mammal a therapeutically effective amount of a compound according to anyone of claims 1 to 27 claim 1 in a unit dosage form.

- 30. (Currently Amended) A method of treating ischemia reperfusion injury in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to anyone of claims 1 to 27 claim 1 in a unit dosage form.
- 31. (Currently Amended) A method of treating myocardial ischemia in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to anyone of claims 1 to 27 claim 1 in a unit dosage form.
- 32. (Currently Amended) A method of treating congestive heart failure in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to anyone of claims 1 to 27 claim 1 in a unit dosage form.
- 33. (Currently Amended) A method of treating arrhythmia in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to anyone of claims 1 to 27 claim 1 in a unit dosage form.
- 34. (Currently Amended) A method of reducing blood clots in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to anyone of claims 1 to 27 claim 1 in a unit dosage form.
- 35. (Currently Amended) A method of treating hypertrophy in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to anyone of claims 1 to 27 claim 1 in a unit dosage form.
- 36. (Currently Amended) A method of treating a disease that arises from thrombotic and

prothrombotic states in which the coagulation cascade is activated in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to anyone of claims 1 to 27 claim 1 in a unit dosage form.

- 37. (Currently Amended) A method of treating diabetes mellitus in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to any one of claims 1 to 27 claim 1 in a unit dosage form.
- 38. (Currently Amended) A method of treating insulin resistance in a mammal comprising concurrently administering to the mammal a therapeutically effective amount of a compound according to anyone of claims 1 to 27 claim 1 in a unit dosage form.
- 39. (Currently Amended) A method of treating hyperinsulinemia in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to any one of claims 1 to 27 in a unit dosage form.
- 40. (Currently Amended) A method of treating diabetes-induced hypertension in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to anyone of claims 1 to 27 claim 1 in a unit dosage form.
- 41. (Currently Amended) A method of treating diabetes-related damage to blood vessels, eyes, kidneys, nerves, autonomic nervous system, skin, connective tissue, or immune system in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to anyone of claims 1 to 27 claim 1 in a unit dosage form.
- 42. (Currently Amended) A method of treating obesity in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to anyone of claims 1 to 27 claim 1 in a unit dosage form.

- 43. (Currently Amended) A compound according to anyone of claims 1 to 27 claim 1 which is a nicotinic acid derivative.
- 44. (Currently Amended) A kit comprising the composition of anyone of claims 1 to 27 claim 1 and instructions for its use in the treatment of a cardiovascular disease, a disease that arises from a thrombotic or prothombotic state in which the coagulation cascade is activated, diabetis, or related diseases.